

4329 FIRST AID KIT - 4329 first aid kit
Honeywell Safety Products USA, INC

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4329 First Aid Kit (1st aid Sp, FABC, EW, aypanal, miralac, amm. Inh, triple, burn spray WS, alcohol wipe, BZK wipes - Z019801)

Eyewash

Active ingredient

Sterile Water 99%

Eyewash

Purpose

Eyewash

Eyewash

Uses

- for flushing the eye to remove loose foreign material, air pollutants or chlorinated water

Eyewash

Warnings

For external use only Obtain immediate medical treatment for all open wounds in or near eyes. To avoid contamination, do not touch tip of container to any surface. Do not reuse. Once opened, discard.

Do not use

- if solution changes color or becomes cloudy
- if you have open wounds in or near the eyes, get medical help right away.

Stop use and ask a doctor if

- you experience eye pain
- changes in vision
- continued redness or irritation of the eye
- condition worsens or persists

Keep out of reach of children

- If swallowed, get medical help or contact a Poison Control Center right away.

Eyewash

Directions

- remove contacts before using
- twist top to remove
- flush the affected area as needed
- control rate of flow by pressure on the bottle
- if necessary, continue flushing with emergency eyewash or shower

Eyewash

Inactive ingredients

sodium chloride, sodium phosphate dibasic, sodium phosphate monobasic

Eyewash

Questions

1-800-430-5490

Ammonia

Active ingredient

Ammonia 15%

Ammonia

Purpose

Respiratory stimulant

Ammonia

Uses

- to prevent or treat fainting

Ammonia

Warnings

For external use only

Do not use

- if you have breathing problems such as asthma or emphysema

Stop use and ask a doctor if

- condition persists

Keep out of reach of children

- If swallowed get medical help or contact a Poison Control Center right away.

Ammonia

Directions

- hold inhalant away from face and crush ampoule between thumb and forefinger at position indicated on sleeve.
- hold near nostrils for inhalation of volatile vapor

Ammonia

Other information

- store at room temperature away from light

Ammonia

Inactive ingredient

alcohol USP, FD&C red #40, lavender oil, lemon oil fcc, nutmeg oil, purified water

Ammonia

Questions or Comments?

1-800-430-5490

Water Soluble 1st Aid Spray

Active ingredient

Benzethonium chloride 0.2% w/w - Benzocaine 10% w/w

Water Soluble 1st Aid Spray

Purpose

Topical antiseptic

Topical anesthetic

Water Soluble 1st Aid Spray

Uses

for temporary relief of pain and itching and helps protect against infection in

- minor cuts and scrapes
- insect bites
- minor skin irritations

Water Soluble 1st Aid Spray

Warnings

For external use only

Flammable

- keep away from fire or flame
- contents under pressure
- do not puncture or incinerate container
- do not expose to temperature above 120 0 F

Do not use

- in the eyes or other mucous membranes
- in cases of serious burns
- in case of deep or puncture wounds
- for a prolonged period of time
- on large portion of the body

Stop use and ask a doctor if

- condition worsens or symptoms persist for more than 7 days
- condition clears up and occurs again within a few days
- redness, swelling, or irritation occurs

Keep out of reach of children

- If swallowed, get medical help or contact a Poison Control Center right away

Water Soluble 1st Aid Spray

Directions

- clean the affected area
- shake can well before using
- hold 4 - 6 inches from surface and spray area until wet
- may be covered with a sterile bandage. If bandaged, let dry first
- for adult institutional use only
- not intended for use on children

Water Soluble 1st Aid pray

Other information

- avoid inhaling
- use only as directed
- intentional misuse by deliberately concentrating and inhaling the contents may be harmful or fatal

Water Soluble 1st Aid pray

Inactive ingredients

dipropylene glycol, isobutane, N-butane, propane

Burn Relief Water Soluble

Active ingredients

Benzethonium chloride 0.2% w/w

Benzocaine 10% w/w

Menthol 0.33% w/w

Burn Relief Water Soluble

Purpose

Topical antiseptic

Topical anesthetic

Topical anesthetic

Burn Relief Water Soluble

Uses

for the temporary relief of pain and itching and helps protect against infection in:

- minor cuts and scrapes
- burns
- sunburn
- insect bites
- minor skin irritations

Burn Relief Water Soluble

Warnings

For external use only

Flammable keep away from fire or flame

- contents under pressure
- do not puncture or incinerate container
- do not expose to temperatures above 120 0 F

Do not use

- in or near the eyes or other mucous membranes
- in case of serious burns
- in case of deep or puncture wounds
- for prolonged period of time
- on large portion of the body

Stop use and ask a doctor if

- condition worsens or symptoms persist for more than 7 days
- condition clears up and recurs within a few days
- redness, swelling, or irritation occurs

Keep out of the reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Burn Relief Water Soluble

Directions

- clean the affected area
- shake can well before using
- hold 4 - 6 inches from surface and spray area until wet
- may be covered with a sterile bandage, if bandaged let dry first
- for adult institutional use only
- not intended for use on children

Burn Relief Water Soluble

Other information

- avoid inhaling
- use only as directed
- intentional misuse by deliberately concentrating or inhaling the contents may be harmful or fatal

Burn Relief Water Soluble

Inactive ingredients

dipropylene glycol, isobutane, n-butane, propane

Triple

Active ingredients

Bacitracin zinc 400 units

Neomycin sulfate (5 mg equivalent to 3.5 mg Neomycin base)

Polymyxin B sulfate 5000 units

Triple Purpose

First aid antibiotic
First aid antibiotic
First aid antibiotic

Triple Uses

- first aid to help prevent infection in
- minor cuts
- scrapes
- burns

Triple Warnings

For external use only

Allergy alert do not use if you are allergic to any of the ingredients

Do not use

- in the eyes
- over large areas of the body

Ask a doctor before use if you have

- a deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- the condition persists or gets worse
- a rash or other allergic reaction develops
- you need to use longer than 1 week

Keep out of the reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Triple Directions

- clean the affected area
- apply a small amount of the product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily
- may be covered with a sterile bandage

Triple Other information

- store at 15 ° to 25 ° C (59 ° to 77 ° F)
- tamper evident sealed packets

- do not use if packet is torn or opened

**Triple
Inactive ingredient**

petrolatum

**Alcohol
Active ingredient**

Isopropyl alcohol 70%

**Alcohol
Purpose**

First aid antiseptic

**Alcohol
Uses**

- first aid to help prevent infection in minor cuts, scrapes, and burns

**Alcohol
Warnings**

For external use only

Flammable, keep away from fire or flame.

Do not use

- in the eyes
- over large areas of the body

Ask a doctor before use if you have

deep or puncture wounds

animal bites

serious burns

When using this product

- do not use longer than one week unless directed by a doctor

Stop use and consult a doctor

- if condition persists or gets worse

Keep out of reach of children

- If swallowed, get medical help or contact a Poison Control Center right away.

**Alcohol
Directions**

- clean the affected area
- apply wipe to affected area 1 to 3 times daily

- may be covered with a sterile bandage
- discard wipe after single use

Alcohol

Other information

store at room temperature 15⁰ to 25⁰ C (59⁰ to 77⁰ F)

Alcohol

Inactive ingredient

water

Miralac

Active ingredient (in each chewable tablet)

Miralac

Active ingredient (in each chewable tablet)

Calcium carbonate 420 mg

Miralac

Purpose

Antacid

Miralac

Uses

for the relief of

- acid indigestion
- heartburn
- sour stomach
- upset stomach associated with these symptoms

Warnings

Ask a doctor before use if you have

- kidney stones
- calcium-restricted diet

Ask a doctor before use if you are

- presently taking a prescription drug. Antacids may interfere with certain prescription drugs

When using this product

- do not take more than 12 tablets in a 24- hour period, or use the maximum dosage of this product for more than 2 weeks, except under the advice and supervision of a doctor.

Keep out of the reach of children.

In case of accidental overdose, seek professional assistance or contact a Poison Control Center immediately.

Miralac

Directions

- chew 1 to 2 tablets every 4 hours as symptoms occur, or as directed by a doctor.

Miralac

Other information

- each tablet contains: calcium 170 mg
- sucrose free
- lactose free
- store at room temperature
- TAMPER EVIDENT PACKETS- DO NOT USE IF OPEN OR TORN

Miralac

Inactive ingredients

magnesium stearate, mint flavor, silicon dioxide, sorbitol, starch

Miralac

Questions or comments?

1-800-430-5490

BZK

Active ingredient

Benzalkonium chloride 0.13% w/v

BZK

Purpose

First aid antiseptic

BZK

Uses

Antiseptic cleansing of face, hands, and body without soap and water

BZK

Warnings

For external use only

Do not use

- in the eyes or over large areas of the body
- on mucous membranes
- on irritated skin
- in case of deep puncture wounds, animal bites or serious burns, consult a doctor
- longer than 1 week unless directed by a doctor

Stop use and ask a doctor if

- if irritation, redness or other symptoms develop
- the condition persists or gets worse

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away

BZK**Directions**

tear open packet and use as a washcloth

BZK**Other information**

- store at room temperature 15⁰ to 30⁰ C (5⁰ - 86⁰ F)
- do not reuse towelette

BZK**Inactive ingredient**

water

BzK**Questions**

1-800-430-5490

Aypanal**Active ingredient (in each tablet)**

Acetaminophen 500 mg

Aypanal**Purpose**

Pain reliever/fever reducer

Aypanal**Uses**

- temporarily relieves minor aches and pains due to the common cold and headache
- temporarily reduces fever

Aypanal**Warnings**

Liver Warning: This product contains acetaminophen. Severe liver damage may occur if you take:

- more than 4,000 mg in 24 hours, which is the maximum daily amount.
- with other drugs containing acetaminophen
- adult has 3 or more alcoholic drinks every day while using this product

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If skin reaction occurs, stop use and seek medical help right away

Do not use

- with any other drug containing acetaminophen (prescription or nonprescription).
- If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.

Ask a doctor before use if you have

- liver disease

Ask a doctor or pharmacist before use if

- you are taking the blood thinning drug warfarin

Stop use and ask a doctor if

- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- new symptoms occur
- redness or swelling is present

If pregnant or breastfeeding

- ask a health professional before use.

Keep out of reach of children.

Keep out of reach of children.

Overdose warning: In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Aypanal

Directions

- **do not take more than directed (see overdose warning)**
- adults and children 12 years of age and over: Take 2 tablets with water every 6 hours while symptoms last.
- do not take any more than 8 tablets in 24 hours.
- children under 12: consult a doctor

Aypanal

Other information

- store at room temperature 15⁰-30⁰C (59⁰-86⁰F)
- TAMPER EVIDENT- DO NOT USE IF OPEN OR TORN

Aypanal

Inactive ingredients

microcrystalline cellulose, povidone, sodium starch glycolate, starch, stearic acid

Aypanal

Questions or Comments

1-800-430-5490

FABC

Active ingredient

Benzalkonium chloride 0.13%

Lidocaine HCl 0.5%

FABC

Purpose

First Aid antiseptic

External analgesic

FABC

Uses

- prevent skin infection
- for temporary relief of pain associated with minor burns

FABC

Warnings

For external use only

Do not use

- in or near the eyes
- if you are allergic to any of the ingredients
- in large areas of the body, particularly over raw surfaces or blistered areas
- for more than 10 days

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- condition worsens
- symptoms persist for more than 7 days or clear up and occurs again within a few days

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

FABC

Directions

- **adults and children 2 years of age and older:**
- clean the affected area

- apply a small amount of this product (equal to the surface area of the tip of a finger) onto affected area 1 to 3 times daily
- may be covered with a sterile bandage
- **children under 2 years of age: consult a doctor**

FABC

Inactive ingredients

aloe barbadensis juice, cetyl alcohol, diazolidinyl urea, edetate disodium, glycerin, glyceryl stearate SE, methylparaben, mineral oil, PEG-100, propylene glycol, propylparaben, stearic acid, trolamine, water

FABC

Questions

1-800-430-5490

4329

Z019801 KIT CONTENTS

1 1X3 PLASTIC 100/BOX
 1 WOVEN 7/8 X 3 50/BOX
 1 FINGERTIP "T" WOVEN 40/BOX
 1 SWIFT KNUCKLE 40/BX
 1 AMMONIA INHALANTS 10 PER
 1 EYE DRESS PKT W/4 ADH STRIPS
 1 TWEEZER PLASTICS 4"
 1 O/H TAPE ADHESIVE TRI-CUT
 1 FIRST AID GUIDE ASHI
 1 BLOODSTOPPER
 10 NON-ADHERENT PADS 2"X3" 10'S
 1 GZE PADS STERILE 3"X 3" 25'S
 2 CTA 3" SINGLE TIP
 1 ANTISEPTIC WIPES BZK CHL 20'S
 1 FIRST AID SPRAY AEROSOL 3 OZ
 1 FIRST AID BURN CREAM 0.9 GRM PKT 20
 1 ALCOHOL WIPES 50'S
 1 AYPANAL EX-STR 2/ENV 250
 1 MIRALAC TABS IND PK 2/ENV 250
 1 BURN SPRAY 3 OZ
 1 TRIPLE BIOTIC .5 GRAM PKT 20
 1 4OZ BFS EYEWASH TRILINGUAL BOTTLE
 1 1 OZ, BUFF EYEWASH

1 SCISSOR BDGE 4" RED PLS HDL
1 400 EMPTY KIT BLANK
1 LBL STOCK 6-3/8"X4"
1 LBL STOCK 4"X2-7/8"
1 LBL CONTS 8"X8",CUSTOM ID B
1 LABEL COVER, GRAINGER Z019801
2 PR LRG NITRILE GLVES
1 SELF-ADH WRAP 2 X 5 YDS NORTH REV D
1 TRI BNDG NON WOVEN 40"X40"X56"
1 COLD PACK UNIT 4"X6" BULK
1 ZIP LOCK BAG 4 X 4" 2 MIL

Eyewash
Principal Display Panel

HoneywellTAMPER-EVIDENT CAP.
TAPA CON SELLO DE SEGURIDAD.
BOUCHON INDICATEUR D'INFRACTION.**eyesaline®****LAVAOJOS
EYESALINE****EYESALINE
EYEWASH****LAVAGE
OCULAIRE
EYESALINE**Solución
Isotónico EstérilSterile
Isotonic SolutionLa Solution
Isotonique Stérile**16 fl. oz. (473 mL)****Drug Facts (for USA only)****Active ingredient** Sterile water 99% **Purpose** Eyewash**Uses**

for flushing the eye to remove loose foreign material, air pollutants, or chlorinated water.

Warnings

For external use only - Obtain immediate medical treatment for all open wounds in or near the eyes. To avoid contamination, do not touch tip of container to any surface. Do not reuse. Once opened, discard.

Do not use

- if solution changes color or becomes cloudy
- if you have open wounds in or near the eyes, get medical help right away

Stop use and consult a doctor if:

- you experience eye pain
- changes in vision
- continued redness or irritation of the eye

condition worsens or persists

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- remove contacts before using
- twist top to remove
- flush the affected area as needed
- control rate of flow by pressure on the bottle
- if necessary, continue flushing with emergency eyewash or shower

Inactive ingredients

sodium chloride, sodium phosphate dibasic, sodium phosphate monobasic

Questions? Call 1-800-430-5490

Honeywell Safety Products USA, Inc. Smithfield, RI. 02917

LABEL #22-204510 Rev. J
REORDER / NUEVO PEDIDO / REAPROVISIONAMIENTO #22-00054-0000

space for lot code and supplier part number

PEEL / PELAR / PEELER

Datos de medicamento (Para EE.UU. solamente)**Ingrediente Activo** Agua estéril 99%**Propósito** Lavados**Usos**

para el lavado de ojo para quitar las partículas sueltas y extrañas, los contaminantes aéros. o agua de cloruro

Advertencias

Para uso externo sólo - Obtenga tratamiento médico inmediato para todas las heridas abiertas en o cerca de los ojos. Para evitar la contaminación, no toque la punta del envase con ninguna superficie. No vuelva a usar. Vez abierto, descarte.

No se usa

- si tiene heridas abiertas en o cerca del ojo, obtenga ayuda médica de inmediato

Deje de usar y consulte a un médico si:

- experimenta dolor de ojo
- cambio de visión
- rojez continuo o irritación del ojo
- la condición empeora o persiste

Manténgase fuera del alcance de los niños.

En caso de ingestión accidental, obtenga atención médica o llame a un Centro de control de envenenamiento inmediatamente.

Instrucciones:

- quite los lentes de contacto antes de usar la solución
- tuerza la tapa para quitar
- enjuague el área afectada según se necesite
- controle el chorro haciendo presión en la botella
- si es necesario, sigue enjuagando con un lavajos o ducha de emergencia

Ingredientes inactivos

cloruro de sodio, fosfato de sodio dibásico, fosfato de sodio monobásico

¿Preguntas? Llame al 1-800-430-5490

Honeywell Safety Products USA, Inc. Smithfield, RI. 02917

Information**Usages**

Pour le rinçage des yeux afin d'enlever un corps étranger, des polluants atmosphériques ou de l'eau chlorée.

Advertissements

Pour usage externe seulement - Obtenir immédiatement des soins médicaux pour toutes les plaies ouvertes dans ou près des yeux. Pour éviter toute contamination, ne pas toucher la pointe du récipient à n'importe quelle surface. Ne pas réutiliser. Une fois ouvert, jetez-les.

Ne pas utiliser

- si la solución a cambiado de color o si ella es brouillée
- si vous avez des plaies ouvertes aux yeux ou à proximité, consultez immédiatement un médecin

Cessez d'utiliser la solution et consulter un médecin

- vous ressentez una dolencia o dolor
- si votre vision change
- rojez o irritación persistante des yeux
- condition empire o persiste

Garder hors de la portée des enfants.

En cas d'ingestion, communiquer immédiatement avec un médecin ou avec un centre antipoison.

Mode d'emploi

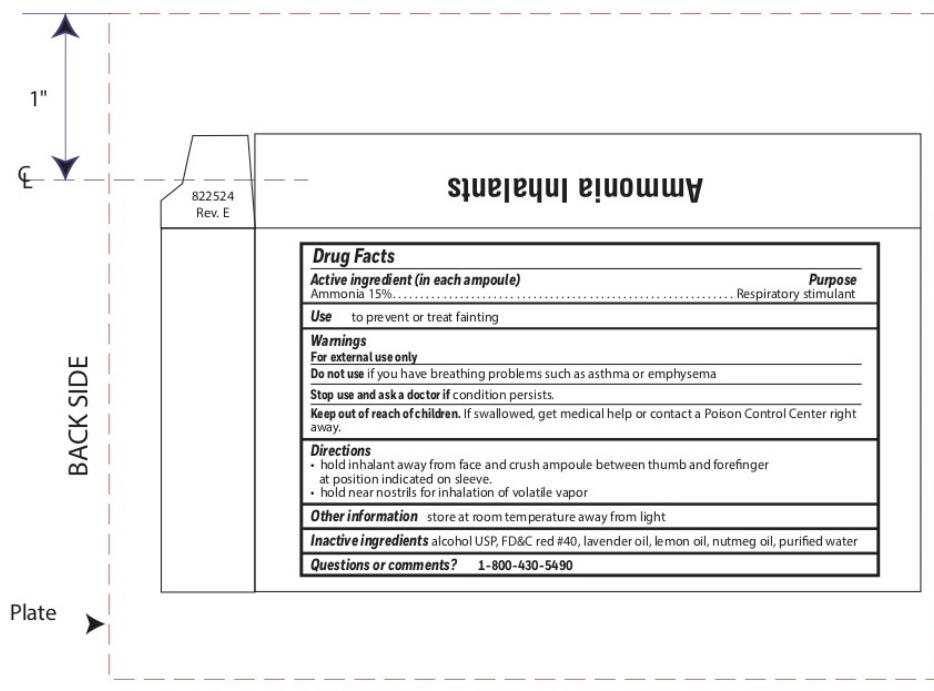
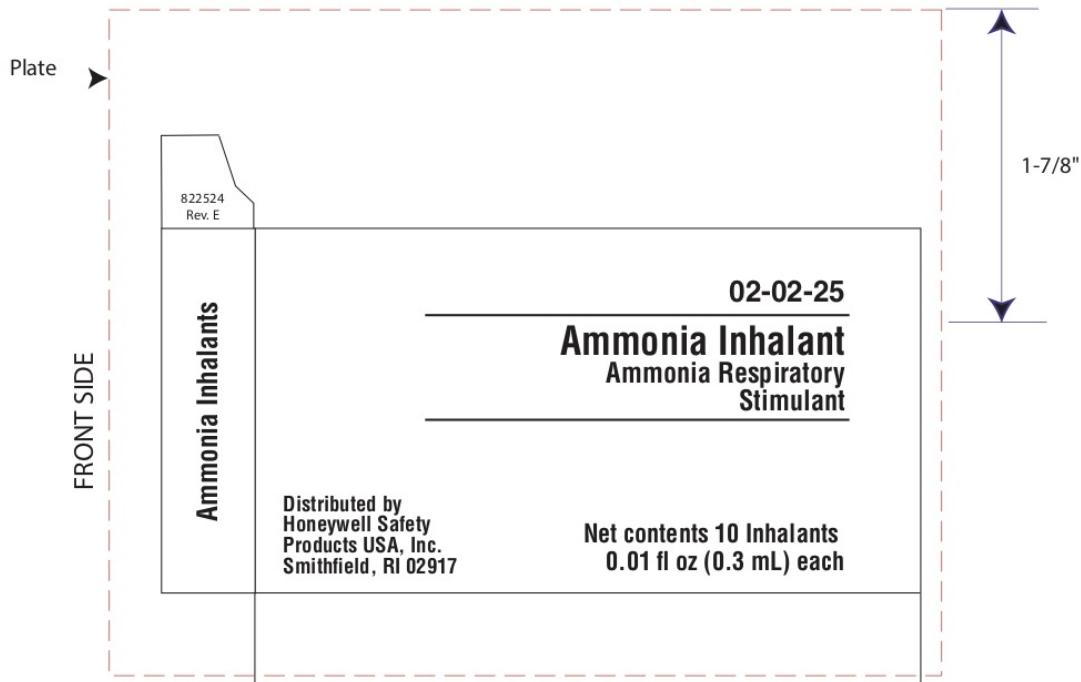
- enlever los lentes de contacto avant l'utilisation
- dévisser le bouchon pour l'enlever
- rincer la zone touchée selon les besoins
- ajuster le débit d'écoulement de la solution en augmentant ou en réduisant la pression exercée sur le contenant
- si nécessaire, continuer de rincer avec unesolution de rinçage oculaire d'urgence ou una douche

Ingredíents agua estéril, cloruro de sodio, fosfato dibásico de sodio, fosfato monobásico de sodio**Des questions?** Faites le 1-800-430-5490

Honeywell Safety Products USA, Inc. Smithfield, RI. 02917

Ammonia
Principal Display Panel

796006 Rev. E Unit Carton Printing Plate for "A" size carton.



796006 Rev. E (page 3 of 3)

Principal Display Panel



47-0303

Cat. No. 151019

DRUG FACTS

Active ingredients

	Purpose
Benzethonium chloride 0.2% w/w.....	Topical antiseptic
Benzocaine 10% w/w.....	Topical anesthetic

Uses

- for the temporary relief of pain and itching and helps to protect against infection in
- minor cuts and scrapes • insect bites • minor skin irritations

Warnings

For external use only

Flammable	• keep away from fire or flame	• contents under pressure
	• do not puncture or incinerate container	• do not expose to temperatures above 120°F

Do not use	• in or near eyes or other mucus membranes	• in case of serious burns
	• in case of deep or puncture wounds	• for a prolonged period of time
	• on large portion of the body	

Stop use and ask a doctor if:

- conditions worsens or symptoms persist for more than 7 days
- condition clears up and recurs within a few days
- redness, swelling or irritation occurs

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- clean the affected area • shake can well before using
- hold 4-6 inches from surface and spray area until wet
- may be covered with a sterile bandage. If bandaged, let dry first
- for adult institutional use only • not intended for use on children

Other information

- avoid inhaling • use only as directed
- intentional misuse by deliberately concentrating and inhaling the contents may be harmful or fatal

Inactive ingredients

dipropylene glycol, isobutane, n-butane, propane

Questions or comments?

1-800-430-5490



Distributed by
Honeywell Safety
Products USA, Inc.
Smithfield, RI 02917

Honeywell

Burn Relief Water Soluble Principal Display Panel



47-0306

Cat. No. 201005

DRUG FACTS

Active ingredients

	Purpose
Benzethonium chloride 0.2% w/w.....	Topical antiseptic
Benzocaine 10% w/w.....	Topical anesthetic
Menthol .33%.....	Topical anesthetic

Uses

- for the temporary relief of pain and itching and helps to protect against infection in
- minor cuts and scrapes • burns • sunburn • insect bites • minor skin irritations

Warnings

For external use only

Flammable	• keep away from fire or flame	• contents under pressure
	• do not puncture or incinerate container	• do not expose to temperatures above 120°F

Do not use	• in or near eyes or other mucus membranes	• in case of serious burns
	• in case of deep or puncture wounds	• for a prolonged period of time
	• on large portion of the body	

Stop use and ask a doctor if:

- conditions worsens or symptoms persist for more than 7 days
- condition clears up and recurs within a few days
- redness, swelling or irritation occurs

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- clean the affected area • shake can well before using
- hold 4-6 inches from surface and spray area until wet
- may be covered with a sterile bandage. If bandaged, let dry first
- for adult institutional use only • not intended for use on children

Other information

- avoid inhaling • use only as directed
- intentional misuse by deliberately concentrating and inhaling the contents may be harmful or fatal

Inactive ingredients

dipropylene glycol, isobutane, n-butane, propane

Questions or comments?

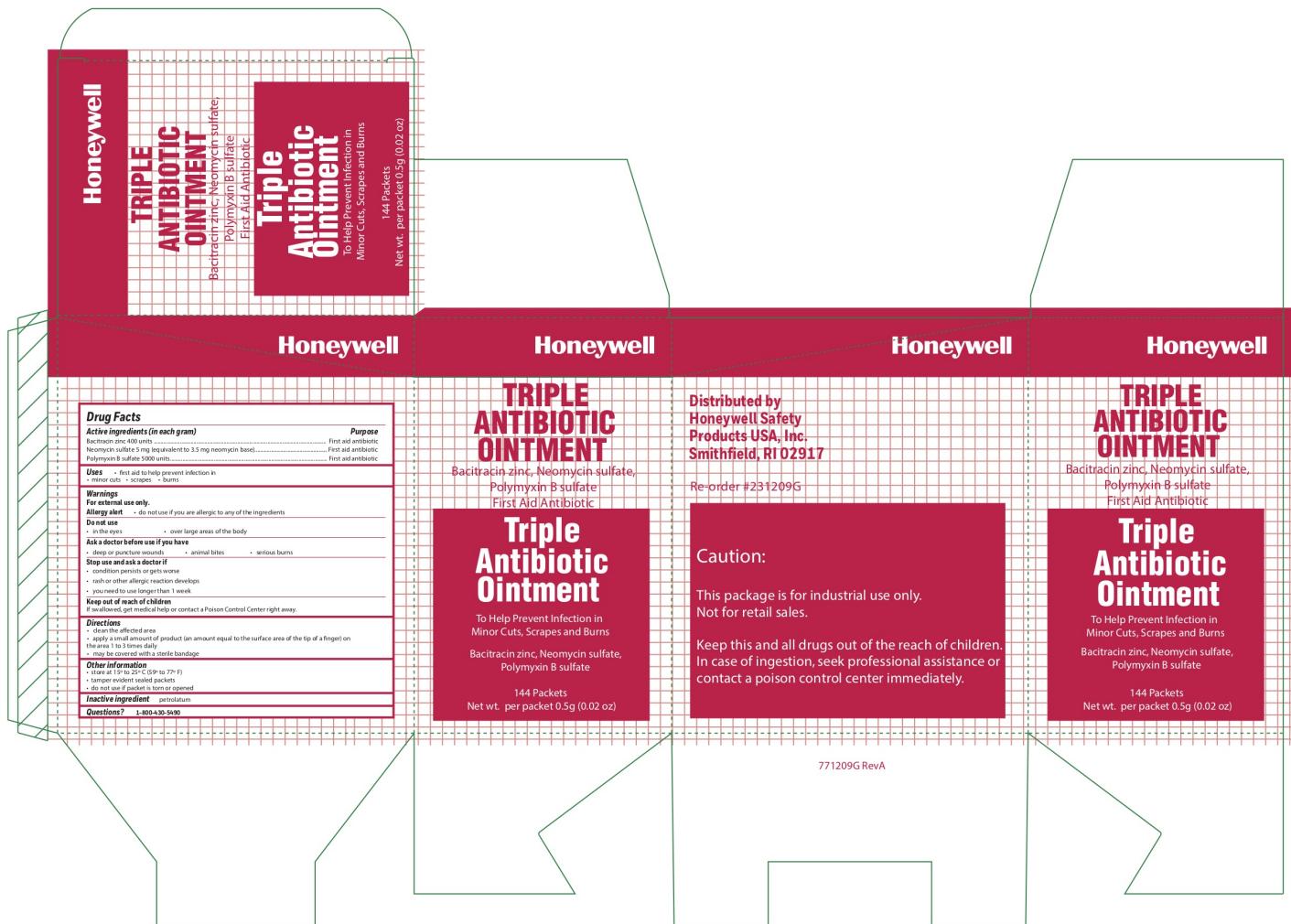
1-800-430-5490



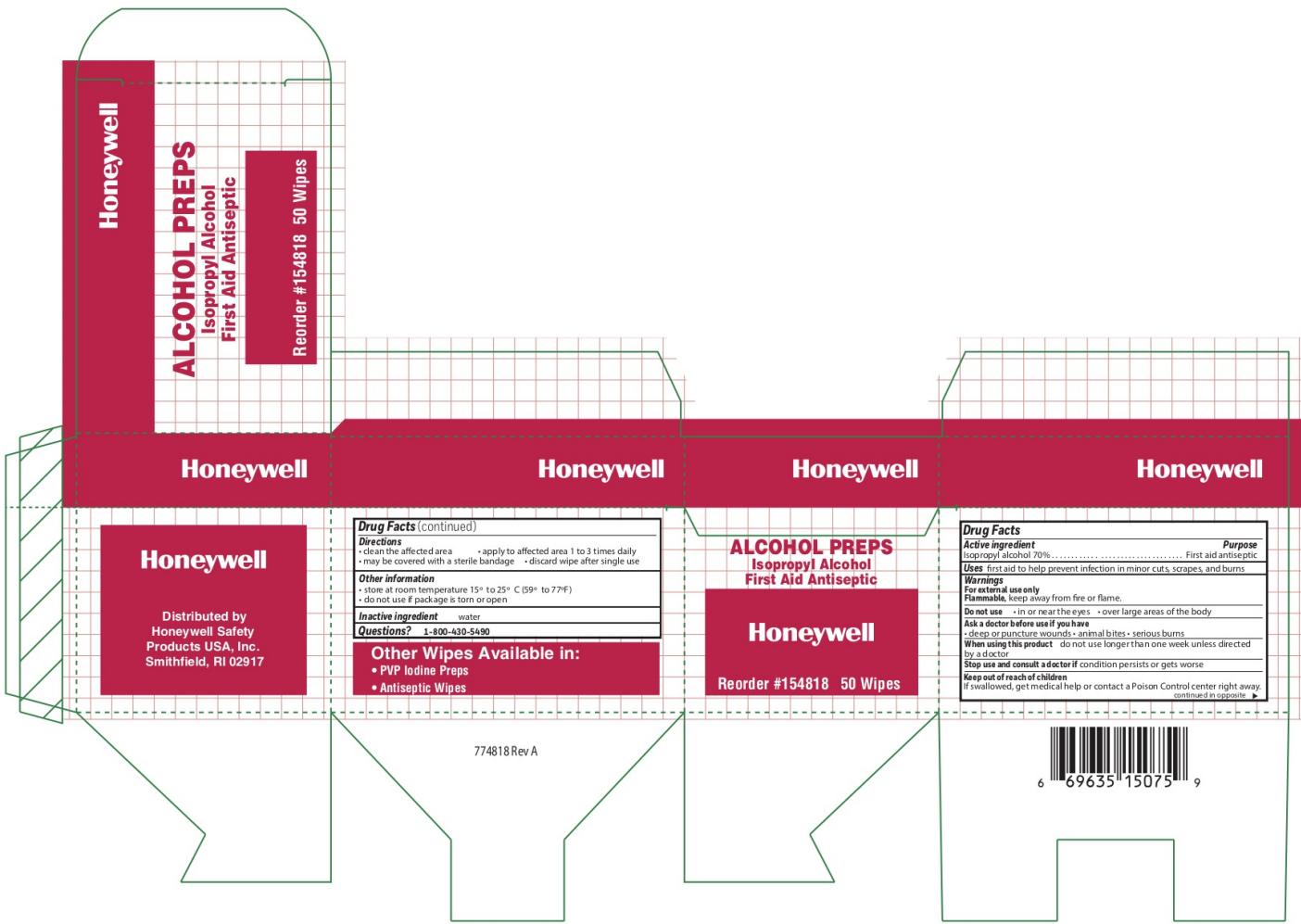
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Honeywell

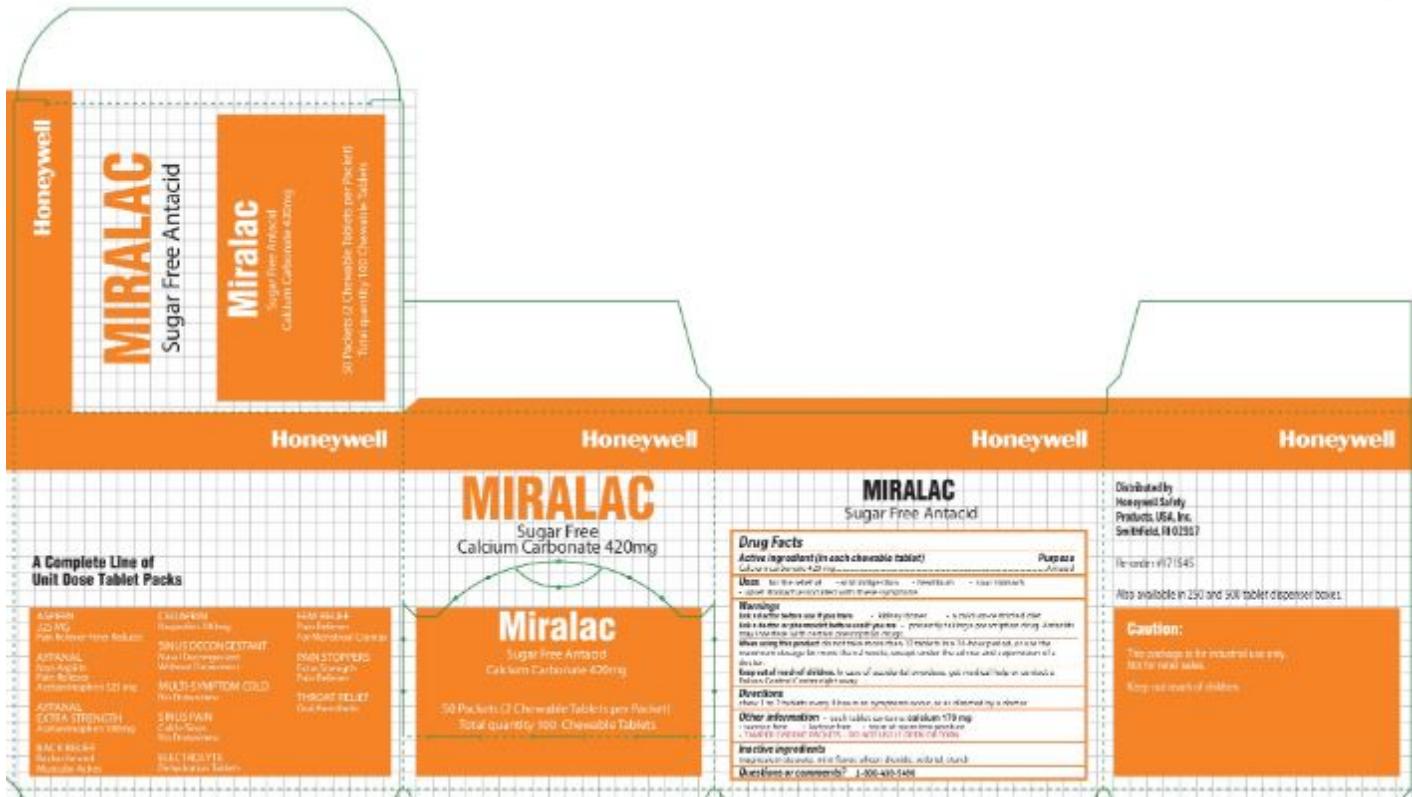
Triple Principal Display Panel



Alcohol Principal Display Panel



Miralac
Principal Display Panel



BZK Principal Display Panel

47001083
Rev B

Antiseptic Towelettes

Honeywell

02-16-35MD

Antiseptic Towelettes

Benzalkonium chloride
First aid antiseptic

Six-Saturated Towelettes

Distributed by
Honeywell Safety
Products USA, Inc.
Smithfield, RI 02917

47001083
Rev B

Antiseptic Towelettes

Honeywell

Drug Facts

Active Ingredient

Benzalkonium chloride 0.133% w/v **Purpose**
First aid antiseptic

Uses

- antiseptic cleaning of face, hands and body without soap and water.
- air dries in seconds

Warnings

For external use only

When using this product • do not use in the eyes or apply over large areas of the body

Ask a doctor before use • in case of deep or puncture wounds, animal bites, or serious burns

Stop use and consult a doctor if

- irritation, redness or other symptoms develop
- condition persists or gets worse

Do not use • longer than 1 week unless directed by doctor

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions • tear open packet, unfold and use as washcloth

Other Information

• store at room temperature 15° -30° C (59° -86° F)

• do not reuse towelette

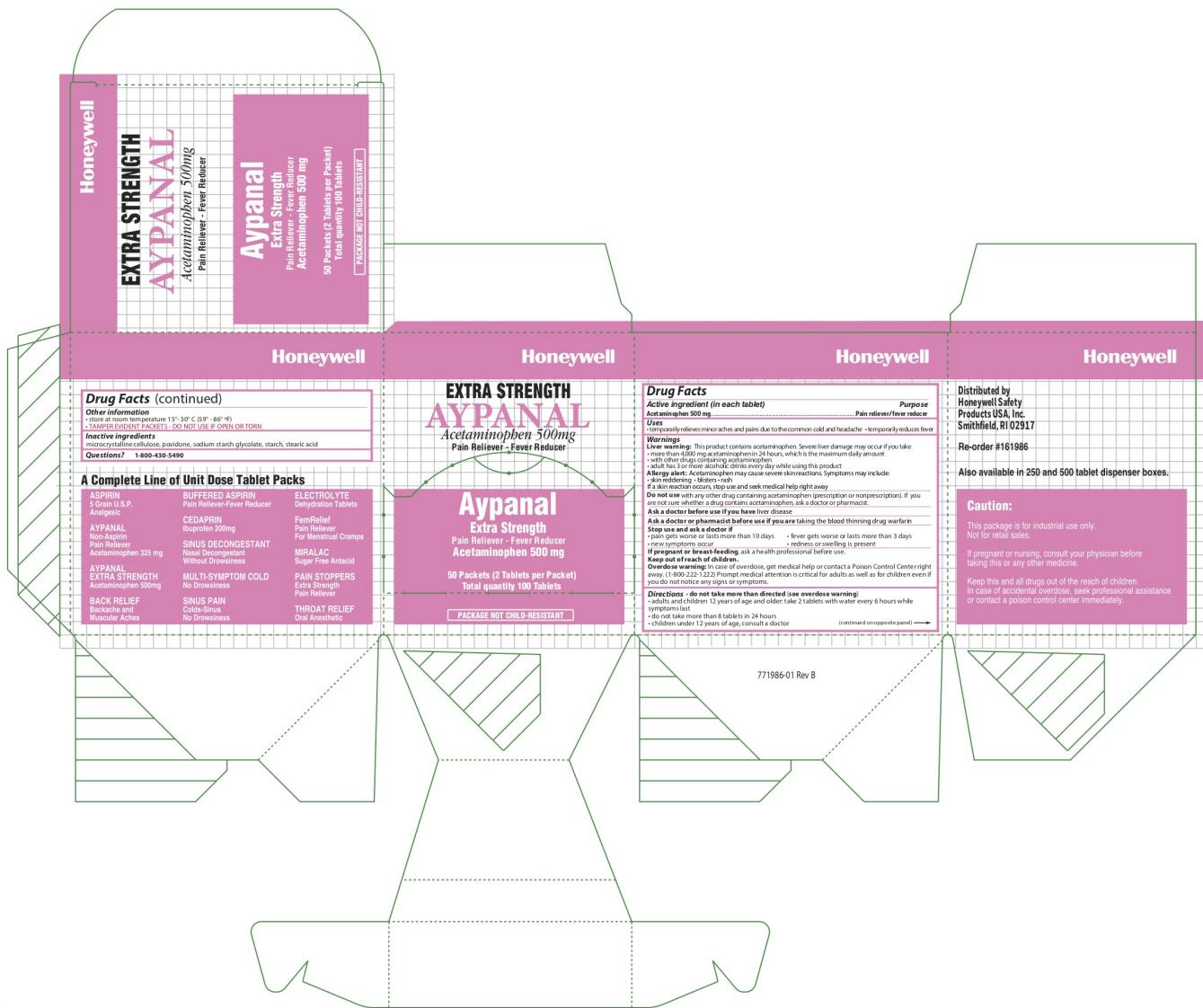
Inactive ingredient

water

Questions or comments

1-800-430-5490

Aypanal
Principal Display Panel



FABC
Principal Display Panel

901850-10

First Aid Burn Cream

021020-10

First Aid Burn Cream

Benzalkonium chloride

First aid antiseptic

Lidocaine HCl

External analgesic

Distributed by
Honeywell Safety
Products USA, Inc.
Smithfield, RI 02917 10 Packets, Net Wt 1/32 oz (0.9g) each

901850-10

First Aid Burn Cream

Drug Facts

Active ingredients

Benzalkonium chloride 0.13%

Lidocaine HCl 0.5%

Purposes

First aid antiseptic

External analgesic

Use - prevent skin infection - for the temporary relief of pain associated with minor burns

Warnings For external use only

Do not use - in or near the eyes - if you are allergic to any of the ingredients
- in large areas of the body particularly over raw surfaces or blistered areas - for more than 10 days

Ask a doctor or before use if you have - deep or puncture wounds - animal bites - serious burns

Stop use and ask a doctor if - condition worsens

- symptoms persist more than 7 days or clear up and occur again within a few days

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

adults and children 2 years and over: - clean the affected area - apply a small amount of this product
equal to the surface area of the tip of a finger onto affected area 1 to 2 times daily

- may be covered with a sterile bandage

children under 2 years: ask a doctor

Other information - tamper evident sealed packets - do not use if packet is opened or torn

Inactive ingredients aloe barbadensis leaf juice, cetyl alcohol, diisodimethyl urea, edetate disodium,
glycerin, glyceryl stearate SE, methyl paraben, mineral oil, PEG-100, propylene glycol, propyl paraben,
stearic acid, triclosan, water

Questions or comments? 1-800-430-5490

**4329 Kit Label
Z019801**

FIRST AID

INDUSTRIAL, BULK
150 PERSON



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Distributed by Honeywell Safety Products USA, Inc. Smithfield, RI 02917

47001713RA

4329 FIRST AID KIT

4329 first aid kit kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0498-4329
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4329-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	148 mL
Part 2	10 AMPULE	3 mL
Part 3	1 CAN	85 g
Part 4	1 CAN	85 g
Part 5	20 PACKET	10 g
Part 6	50 POUCH	20 mL

Part 7	125 PACKET	250
Part 8	20 PACKET	28 mL
Part 9	125 PACKET	250
Part 10	20 PACKET	18 g

Part 1 of 10

EYESALINE EMERGENCY EYEWASH

purified water liquid

Product Information

Item Code (Source)	NDC:0498-0100
Route of Administration	OPHTHALMIC

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	98.6 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE (UNII: 593YOG76RN)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0100-02	118 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:0498-0100-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	12/18/2018	

Part 2 of 10

AMMONIA INHALENT

ammonia inhalent inhalant

Product Information

Item Code (Source)	NDC:0498-3334
Route of Administration	RESPIRATORY (INHALATION)

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AMMONIA (UNII: 5138Q19F1X) (AMMONIA - UNII:5138Q19F1X)	AMMONIA	0.045 g in 0.3 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-3334-00	0.3 mL in 1 AMPULE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/18/2018	

Part 3 of 10

FIRST AID ANTISEPTIC WATER SOLUBLE

benzethonium chloride, benzocaine spray

Product Information

Item Code (Source)	NDC:0498-0031
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.2 g in 100 g
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	10 g in 100 g

Inactive Ingredients

Ingredient Name	Strength

ISOBUTANE (UNII: BXR49TP611)

BUTANE (UNII: 6LV4FOR43R)

PROPANE (UNII: T75W9911L6)

DIPROPYLENE GLYCOL (UNII: E107L85C40)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0031-40	85 g in 1 CAN; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	09/19/2018	

Part 4 of 10

BURN WATER SOLUBLE

benzocaine, benzethonium chloride, menthol spray

Product Information

Item Code (Source)	NDC:0498-0021
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.2 g in 100 g
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	10 g in 100 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.33 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ISOBUTANE (UNII: BXR49TP611)	
BUTANE (UNII: 6LV4FOR43R)	
PROPANE (UNII: T75W9911L6)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0021-40	85 g in 1 CAN; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	11/12/2018	

Part 5 of 10

TRIPLE ANTIBIOTIC

bacitracin zinc, polymyxin b sulfate, neomycin sulfate ointment

Product Information

Item Code (Source)	NDC:0498-0750
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [iU] in 1 g
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

Product Characteristics

Color	white	Score
Shape		Size
Flavor		Imprint Code
Contains		

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0750-36	0.5 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	09/19/2018	

Part 6 of 10

ALCOHOL WIPE

isopropyl alcohol swab

Product Information

Item Code (Source)	NDC:0498-0143
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0143-04	0.4 mL in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/18/2018	

Part 7 of 10

MIRALAC

calcium carbonate tablet

Product Information

Item Code (Source)	NDC:0498-0303
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6ZB, CARBONATE ION - UNII:7UJQ5OPE7D)	CALCIUM CARBONATE	420 mg

Inactive Ingredients

Ingredient Name	Strength
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SORBITOL (UNII: 506T60A25R)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	white	Score	2 pieces
Shape	ROUND	Size	11mm
Flavor	MINT	Imprint Code	FR8
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		2 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part331	02/22/2012	

Part 8 of 10

ANTISEPTIC TOWELETTE

benzalkonium chloride liquid

Product Information

Item Code (Source)	NDC:0498-0501
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name		Strength
WATER (UNII: 059QF0KO0R)		

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0501-00	1.4 mL in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	12/22/2017	

Part 9 of 10

AYPANAL EX

acetaminophen tablet

Product Information

Item Code (Source)	NDC:0498-2110
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINO PHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg

Inactive Ingredients

Ingredient Name	Strength
MICRO CRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
POVIDONE (UNII: FZ989GH94E)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	12mm
Flavor		Imprint Code	FR1
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-2110-01	2 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	01/02/2017	

Part 10 of 10

FIRST AID BURN

benzalkonium chloride, lidocaine hydrochloride cream

Product Information

Item Code (Source)	NDC:0498-0903	
Route of Administration	TOPICAL	

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
METHYLPARABEN (UNII: A2I8C7H9T)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
PEG-100 STEARATE (UNII: YD01N1999R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
EDETA DISODIUM (UNII: 7FLD91C86K)	
TROLAMINE (UNII: 9O3K93S3TK)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
DIAZOLIDINYLMUREA (UNII: H5RIZ3MPW4)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		0.9 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	12/20/2017		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		10/18/2018		

Labeler - Honeywell Safety Products USA, INC (079287321)

Establishment				
Name	Address	ID/FEI	Business Operations	
James Alexander		040756421	manufacture(0498-3334)	

Establishment				
Name	Address	ID/FEI	Business Operations	
Honeywell Safety Products USA, INC		079287321	pack(0498-4329)	

Establishment				
Name	Address	ID/FEI	Business Operations	
Ultra Seal Corporation		085752004	manufacture(0498-2110, 0498-0303)	

Establishment				
Name	Address	ID/FEI	Business Operations	
Dixon Investments		115315822	manufacture(0498-0031, 0498-0021)	

Establishment				
Name	Address	ID/FEI	Business Operations	
Water-Jel Technologies		155522589	manufacture(0498-0750, 0498-0903)	

Establishment				
Name	Address	ID/FEI	Business Operations	
Honeywell Safety Products USA, Inc.		167518617	manufacture(0498-0100)	

Establishment				
Name	Address	ID/FEI	Business Operations	
Changzhou Maokang Medical		421317073	manufacture(0498-0143, 0498-0501)	

Revised: 5/2019

Honeywell Safety Products USA, INC